AMENDMENTS TO THE CLAIMS

Please amend claims 1, 8, 9, 15, 23, 24, 28, 29 and 31 and cancel claims 3, 4, and 30.

1. (Currently Amended) A method of forming a semiconductor device having a metal silicide by a low pressure chemical vapor deposition (LPCVD) process using a source gas of TiCl₄, comprising the steps of:

forming a source/drain junction area on a silicon substrate;

forming an attack protection layer on the source/drain junction area, wherein the attack protection layer is electrically conductive and prevents a silicon substrate attack caused by chlorine (Cl) gas gases generated from the source gas of TiCl₄;

forming a titanium (Ti) layer over the attack protection layer through a low pressure chemical vapor deposition (LPCVD) the LPCVD process using a source gas of TiCl₄; and

illuminating an ultra violet light having a higher energy than a binding energy of a

SiCl reaction product on the surface of the Ti layer to remove remnant chlorine (Cl)

components in the Ti layer; and

diffusing the Ti layer into the attack protection layer to thereby form a metal silicide layer.

- 2. (Original) The method as recited in claim 1, wherein a poly-silicon layer formed by using a chemical vapor deposition (CVD) process is used for forming the attack protection layer.
 - 3. (Canceled)
 - 4. (Canceled)

5. (Original) The method as recited in claim 2, wherein the attack protection layer is formed by the CVD process using a source gas of Si₂H₆/Cl/H₂.

- 6. (Original) The method as recited in claim 2, wherein a thickness of the attack protection layer ranges from about 50 Å to about 200 Å
- 7. (Original) The method as recited in claim 5, wherein the CVD process for forming the attack protection layer is carried out at a temperature ranging from about 600 °C to about 700 °C and at a pressure ranging from about 0.1 mtorr to about 1.0 mtorr.
- 8. (Currently Amended) The method as recited in claim 5, further comprising the step of deoxidizing the surface of the attack protection layer by using hydrogen (H₂) gas to remove the remnant chlorine (Cl) radical components in the attack protection layer after depositing the attack protection layer.
- 9. (Currently Amended) The method as recited in claim 5, further comprising the step of illuminating an ultra violet light having a higher energy than a binding energy of SiCl on the surface of the attack protection layer to remove the remnant chlorine (Cl) radical components in the attack protection layer[[;]].
- 10. (Original) The method as recited in claim 1, wherein a titanium nitride (TiN) layer formed by using a chemical vapor deposition (CVD) process is used for forming the attack protection layer.
- 11. (Original) The method as recited in claim 10, wherein the TiN layer is deposited by using the TiCl₄ source gas added with ammonia (NH₃) gas in an identical chamber where the Ti layer is subsequently deposited.

12. (Original) The method as recited in claim 10, wherein a thickness of the attack protection layer ranges from about 50 Å to about 200 Å.

- 13. (Original) The method as recited in claim 1, wherein the Ti layer is deposited by using the LPCVD process at a temperature ranging from about 600 °C to about 700 °C and at a pressure ranging from about 1 torr to about 50 torr.
- 14. (Original) The method as recited in claim 13, wherein the LPCVD process is performed by using the TiCl₄ source gas added with ammonia (NH₃) gas and hydrogen (H₂) gas including argon (Ar) gas and a flow amount ratio of the NH₃ gas to the Ar gas is about 1 to about 5.
- 15. (Currently Amended) A method for forming a barrier metal layer for a semiconductor device fabrication, comprising the steps of:
- a) forming a contact hole exposing an active area through a selective etch of an insulation layer formed on a silicon substrate providing the active area;
- b) forming an attack protection layer for preventing the silicon substrate attack caused by a succeeding titanium layer deposition process on the active area exposed by the contact hole, wherein the attack protection layer is electrically conductive;
- c) forming a titanium (Ti) layer along a profile of the attack protection layer formed on the active area by using a low pressure chemical vapor deposition (LPCVD) process using a source gas of TiCl₄;
- d) diffusing the Ti layer into the attack protection layer, to thereby forming a metal silicide layer; and

e) illuminating an ultra violet light having a bigger energy than a binding energy of SiCl on the surface of the Ti layer to remove remnant chlorine (Cl) components in the Ti layer; and

- e) f) forming a titanium nitride (TiN) layer on the Ti layer.
- 16. (Original) The method as recited in claim 15, wherein a poly-silicon layer formed by a chemical vapor deposition (CVD) process is used as the attack protection layer.
- 17. (Original) The method as recited in claim 16, wherein the CVD process is carried out by using a source gas of Si₂H₆/Cl/H₂.
- 18. (Original) The method as recited in claim 17, wherein the CVD process for forming the attack protection layer is performed at a temperature ranging from about 600 °C to about 700 °C and at a pressure ranging from about 0.1 mtorr to about 1 mtorr.
- 19. (Original) The method as recited in claim 15, wherein a titanium nitride (TiN) layer formed by a chemical vapor deposition (CVD) process is used as the attack protection layer.
- 20. (Original) The method as recited in claim 19, wherein the attack protection layer is deposited by using the TiCl₄ source gas added with ammonia (NH₃) gas in an identical chamber where the Ti layer will be deposited.
- 21. (Original) The method as recited in claim 19, wherein a thickness of the attack protection layer ranges from about 10 Å to about 100 Å

22. (Original) The method as recited in claim 15, wherein the Ti layer is deposited by using the LPCVD process at a temperature ranging from about 600 °C to about 700 °C and at a pressure ranging from about 1 torr to about 50 torr.

- 23. (Currently Amended) The method as recited in claim 22, wherein the LPCVD process is performed by using the TiCl₄ source gas added with ammonia (NH₃) gas and hydrogen (H₂) gas including argon (Ar) gas and a flow amount ratio of the NH₃ gas to the Ar gas is about 1 to about $5\frac{1}{5}$.
 - 24. (Currently Amended) The method as recited in claim 15, wherein the TiN layer is deposited on the Ti layer by using a low pressure chemical vapor deposition (LPCVD) the LPCVD process at a temperature ranging from about 600 °C to about 700 °C and at a pressure ranging from about 1 torr to about 50 torr.
- 25. (Original) The method as recited in claim 24, wherein the LPCVD process is performed by using the TiCl₄ source gas added with ammonia (NH₃) gas and hydrogen (H₂) gas including argon (Ar) gas and a flow amount ratio of the NH₃ gas to the Ar gas is about 8 to about 15.
- 26. (Original) The method as recited in claim 25, wherein the TiN layer is deposited in an identical chamber where the Ti layer is deposited.
- 27. (Original) The method as recited in claim 15, wherein the titanium silicide layer is produced by carrying out a heat treatment process at a temperature ranging from about 700 °C to about 900 °C.

28. (Currently Amended) The method as recited in claim 15, wherein further comprising the step of deoxidizing the surface of the Ti layer using hydrogen (H₂) gas to remove a remnant chlorine radical components after depositing the Ti layer and the TiN layer.

- 29. (Currently Amended) The method as recited in claim 17, further comprising the step of deoxidizing the surface of the attack protection layer using a hydrogen (H₂) gas to remove remnant chlorine radical components after depositing the attack protection layer, i.e., the poly-silicon layer.
 - 30. (Canceled)
- 31. (Currently Amended) The method as recited in claim 17, further comprising the step of illuminating an ultra violet light having a bigger energy than a binding energy of SiCl to remove remnant chlorine (Cl) radical components in the attack protection layer.